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Ambient formaldehyde measurements made at a remote marine boundary layer site during the NAMBLEX campaign – a comparison of data from chromatographic and modified Hantzsch techniques

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Abstract

Ambient formaldehyde concentrations are reported from the North Atlantic Marine Boundary Layer Experiment (NAMBLEX) campaign at Mace Head on the west coast of Ireland during August 2002. The results from two techniques, using direct determination via gas chromatography and the Hantzsch technique, show similar trends but a significant off set in concentrations. For westerly air flows characteristic of the marine boundary layer, formaldehyde concentrations from the gas chromatographic and Hantzsch technique ranged from 0.78–1.15 ppb and 0.13–0.43 ppb, respectively. Possible reasons for the discrepancy have been investigated and are discussed, however, no satisfactory explanation has yet been found. In a subsequent intercomparison the two techniques were in good agreement.

The observed concentrations have been compared with previous formaldehyde measurements in the North Atlantic marine boundary layer and with other measurements from the NAMBLEX campaign. The measurements from the Hantzsch technique and the GC results lie at the lower and upper ends respectively of previous measurements. In contrast to some previous measurements, both techniques show distinct diurnal profiles with day maxima and with an amplitude of approximately 0.15 ppb. Strong correlations were observed with ethanal concentrations measured during NAMBLEX and the ratio of ethanal to formaldehyde determined by the gas chromatographic technique is in good agreement with previous measurements.

Some simple box modelling has been undertaken to investigate possible sources of formaldehyde. Such models are not able to predict absolute formaldehyde concentrations as they do not include transport processes, but the results show that oxygenated VOCs such as ethanal and methanol are very significant sources of formaldehyde in the air masses reaching Mace Head.

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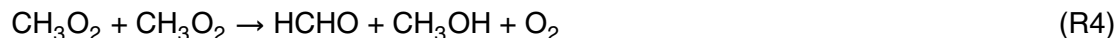
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1. Introduction

The North Atlantic Boundary Layer Experiment, NAMBLEX, took place at Mace Head, Eire, during July and August 2002 to help quantify our understanding of photochemical oxidation processes in clean and moderately polluted environments. Objectives included quantifying the role of halogen species in the marine boundary layer (MBL), studying the reactive nitrogen budget and formation of new particles (Read et al., 2005). Objectives particularly relevant for our formaldehyde (HCHO), measurements included model/measurement comparisons for radical species and the role of reactive hydrocarbons in the MBL. Formaldehyde plays an important role in determining radical concentrations, influencing both HO_x formation and removal. Due to its importance and difficulties in determining accurate concentrations (Gilpin et al., 1997) formaldehyde concentrations were measured using two different techniques. The University of Leeds (UoL) used a chromatographic method based on the detection of separated HCHO with a helium ionization detector (Hunter et al., 1998, 1999; Hopkins et al., 2003). The University of East Anglia, UEA, used a version of the Hantzsch reaction (Cardenas et al., 2000) where ambient HCHO was scrubbed into solutions and derivitized by reaction with 2,4 pentadione and ammonia. The adduct was then detected via fluorescence at 510 nm following UV excitation by a mercury lamp. The two systems were located at sites approximately 200 m apart.

In the very remote environment, methane is the major source of formaldehyde via the reactions:



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5 Under such low NO_x conditions, the methyl peroxy radical (CH_3O_2) formed from the reaction of OH with CH_4 predominantly reacts with other peroxy radicals. Reaction with HO_2 leads to CH_3OOH which effectively acts as a reservoir species on route to HCHO formation (although it can be rained out). Self reaction leads more directly to HCHO formation. Reaction (8) is very rapid and other reactions of methoxy radicals do not need
10 to be considered. Additional sources of formaldehyde include higher hydrocarbons and oxygenated VOCs such as methanol or acetaldehyde.

In the presence of sufficient NO, CH_3O_2 radicals react directly with NO to form methoxy radicals and subsequently formaldehyde via Reaction (R8).



15 In a modelling study of formaldehyde production in the remote Indian Ocean, Wagner et al. (2002) showed that in the presence of only 2 pptv of NO approximately 50% of CH_3O_2 reacts via Reaction (9).

There are two major chemical removal processes for formaldehyde. Firstly, reaction with OH (10) which, via the subsequent rapid reaction of HCO with oxygen (11), is a
20 major route for conversion of OH to HO_2 .



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Secondly, photolysis, which can act as a significant free radical source. The efficiency of HCHO as a radical source depends on the branching ratio between the molecular and radical channels (Pope et al., 2005):



Formaldehyde, and carbonyls in general, have been shown to be major sources of HO_x radicals in the urban winter atmosphere (Heard et al., 2004), but formaldehyde is also an important HO₂ source in the remote free troposphere. Frost et al. (2002) noted that the importance of formaldehyde as a radical source will increase at higher, increasingly drier altitudes.

10 Table 1 lists some previous formaldehyde measurements in remote MBL environments, focusing particularly on studies in the North Atlantic environment. Measurements have been with a variety of techniques and from airborne, shipborne and coastal platforms. Average values range from ~200–1000 pptv. A seasonal dependence has been observed for formaldehyde measurements in the EMEP programme (Solberg et al., 1996) peaking during the summer months. There is conflicting evidence on meridional variations with measurements in the North Atlantic showing both positive and negative variations with increasing latitude (Harris et al., 1992; Weller et al., 2000). Fried and co-workers have carried out several airborne campaigns, their measurements show a general decrease in formaldehyde concentrations with altitude (Fried et al., 2002, 2003).

25 As can be seen from Table 1, a variety of experimental techniques have been deployed. Partially, this reflects the difficulties in making reliable measurements on this important atmospheric intermediate. The deployment of a variety of techniques has highlighted potential systematic errors in the measurement techniques. A good review of several techniques and the results of a field intercomparison can be found in the work by Gilpin et al. (1997).

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2. Experimental

2.1. Site

NAMBLEX took place at the Mace Head observatory on the west coast of Eire (53°19'34" N, 9°54'14" W) during July and August 2002. A majority of formaldehyde measurements were taken between 1–21 August 2002. The location of the site is shown in Fig. 1. Air arriving between the angles of 180 and 300° is free from any local land influence.

Five day back trajectories were calculated based on the wind field analysis produced by the European Centre of Medium Range Weather Forecasts (ECMWF) and were used to classify the origins of the air masses arriving at Mace Head. During the campaign local meteorological measurements were made by a wind profiler and its observations provided a record of the boundary layer structure, including measurements of average wind speed and direction (Norton et al., 2005). The local measurements suggest that for westerly winds, the ECMWF trajectories and local measurements are in good agreement. For north-easterly wind flows, characteristic of the early part of the campaign, local sea breezes dominated. However, the chemical signatures of typical anthropogenic pollutants such as CO and C₂H₂ suggest that the site is still receiving air broadly characteristic of origin of the air mass, albeit potentially moderated by local meteorology.

The inset to Fig. 1 shows the location of the two formaldehyde measurements. The UEA apparatus was co-located with a majority of the other instrumentation ~100 m distant and at ~10 m elevation from the average high tide mark. The UoL instrumentation was located a further ~200 m inshore and elevated by a further ~20 m.

2.2. UoL apparatus

The UoL instrument used during the NAMBLEX campaign was based on a gas chromatographic (GC) system as described by Hopkins et al. (2003). The sampling inlet

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was placed 2 m above the ground and consisted of ~12 m 1/4" PFA tubing, and was pumped at a speed of 1 slm. The sample passed through a 6.4 ml sample loop (Silco Steel). During injection, helium carrier gas (CP grade, BOC, 9 ml min⁻¹, backing pressure 50 psi, further purified via passing through liquid nitrogen traps) was diverted through the loop, sweeping the sample onto the column (50 m, 0.32 mm id, 100% dimethyl polysiloxane, WCOT column, 5 μm phase thickness, CP-Sil 5CB Chrompack, Netherlands) and refocused at the head of the column in a liquid nitrogen trap. Following the elution of the untrapped air, the analytes were released and were separated in the column and detected using an argon doped (1% Ar in He, BOC, CP grade), pulsed discharge helium ionisation detector (pdHID) (Model D4, VICI AG, Schenkon, Switzerland). The detector flow was 30 ml min⁻¹ maintained with a calibrated restrictor. After the elution of formaldehyde the column flow was reversed and the column was back flushed (30 ml min⁻¹, 70 s, backing pressure 60 psi) to prevent water reaching the detector or the build-up of heavier weight material on the column. The sample cycle time throughout the campaign was 5.5 min.

The detection limit during the campaign was 42 pptv; this was calculated taking the minimum detectable peak to have a signal to noise ratio of 3:1. The system used a formaldehyde gas phase standards generator for calibration (KIN-TEK, LaMarque, TX). A permeation tube containing polymeric formaldehyde with a known emission rate at a set reference temperature (333 K) was held in a stabilized oven (temperature variation <±0.1 K) flushed with a constant flow rate (10.0 sccm) of UHP nitrogen. Delivery concentrations of 4–50 ppbv (during campaign), or subsequently 2–50 ppbv (with increased flow rate), were obtained by diluting the oven flow with additional known flow rates of UHP nitrogen. The delivery tube was connected to the GC sample inlet via a 'tee' junction with a positive flow out of the vent, ensuring that the delivery pressure was at one atmosphere. The precision was attained from the standard deviation of replicate calibration factors and was calculated to be 3%. The uncertainty was attained from the errors associated with the standards generator and was determined to be 9% (2σ).

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2.3. UEA apparatus

The fluorescence technique is based on the Hantzsch reaction which is a liquid phase reaction of formaldehyde followed by fluorescence detection of the resulting adduct. This technique requires formaldehyde to be transferred from the gas phase into liquid phase, achieved via a stripping solution of 0.1 N H₂SO₄ (made up from ACS reagent grade H₂SO₄, Aldrich) at room temperature, which is pumped through a coil (45 cm long, 10 turns of 3 mm o.d. Pyrex) and forced into contact with gaseous formaldehyde. Ambient formaldehyde was sampled at a height of ~5 m through approximately 10 m of 3/8" PFA tubing. The pumping speed was a total of 15 slm giving a very short residence time in the sampling line, and only small decrease in pressure in the sample line. The sample was introduced into the instrument at 1.5 slm through a 1 m length of 1/4" PFA tubing.

The Hantzsch reaction is based on the liquid phase reaction of formaldehyde with acetylacetone (2,4 pentadione) and ammonia to produce diacetyldihydrolutidine, (DDL) that is excited at 412 nm (Hg-Phosphor 215 lamp, Jelight, USA) and the fluorescence is detected at 510 nm with a photomultiplier tube (Hamamatsu).

The detection limit for this instrument was in the region of 50 pptv (Cardenas et al., 2000). The instrument was also calibrated using a permeation source (KIN-TEK LaMarque, TX). The calibrant introduced into the instrument via a "Tee-piece" to vent the excess and ensure the calibration was delivered at atmospheric pressure. The precision of the instrument was ±6 % and the accuracy ±13% with a minimum error of ±25 pptv. Zeros were performed every 5 h by scrubbing the sample with a charcoal filter.

2.4. Intercomparisons and artefacts

During the campaign the calibration sources for both instruments were exchanged and compared. When used on the UoL instrument, the UEA calibration source produced values within 2% of those generated using the UoL source. It was therefore concluded

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that the calibration sources were in good agreement for the campaign period.

The two instruments also took part in a laboratory intercomparison at the National Physics Laboratories (Teddington, UK) during July 2003. In these experiments all the instruments sampled from a common chamber (NPL Standard Atmosphere Generator). HCHO concentrations in the chamber were calculated from permeation rates of a known standard and measured dilution flows. All the instruments performed linearity and stability checks and a blind sampling. Various concentrations of ozone were generated and introduced into the chamber to investigate possible artefacts.

Both the instruments responded well to the linearity and stability experiments and reported results within experimental errors for the blind testing, although the concentration, at ~ 7 ppbv, was significantly higher than ambient concentrations in the MBL. Both instruments showed a positive interference for ozone as shown in Fig. 2.

The magnitude of the ozone artefact on the UoL instrument was significantly greater, but was reproducible, both at NPL and during subsequent experiments at Leeds. The results reported below for both instruments have been corrected for ozone measurements as recorded at Mace Head by the FAGE group using a UV photometric analyser.

3. Results

3.1. Comparison of time series measurements from UoL and UEA apparatus

Figure 3 shows the formaldehyde time series for both instruments. Two observations are immediately apparent. Firstly, the results from the UoL are significantly higher, and outside the combined random errors of both techniques (9% for UoL and 13% for UEA based on errors associated with the calibration processes). Secondly there is a reasonable correlation between the two measurements, and in some cases, for example that shown in the inset to Fig. 3, the pattern of variation is almost identical.

Figure 4 shows a correlation plot of the results. There is significant scatter in the data which may be associated with sampling slightly different air masses either due

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to local meteorology or to differences in the sampling techniques. The UoL apparatus is essentially a grab sample taken every 5.5 min, whereas UEA data are based on averaging over a one minute period. The difference in the UoL and UEA data appears to be in an offset rather than in the gradient of the correlation plot.

The two most obvious sources of such an offset would be a significant blank or interference in the UoL apparatus or loss of sample in the UEA apparatus. Blanks were recorded on a daily basis for the UoL apparatus. These were obtained by attaching a cylinder of zero nitrogen (Premier Grade, Air Products) via $\frac{1}{4}$ " Teflon tubing to the inlet port of the GC. A 'tee' junction was placed upstream of the sample inlet to restrict the upstream pressure to 1 atmosphere. The zero air was drawn into the GC via the pump in the normal way. An ozone artefact for the UoL apparatus has already been described. Corrections were typically of the order of 300–400 pptv formaldehyde. We cannot rule out the possibility of other undetected artefacts.

Significant loss processes in the UEA instrument would be very surprising since the whole wetted path of the sample is PFA and no loss processes on PFA have been seen in numerous laboratory tests. The difference in pressure between the sample drawn down the $\frac{3}{8}$ " PFA and the calibrant delivered directly into the instrument is at most 10%. Both losses and differences in pressure would lead to an underestimation of HCHO, but would be expected to show a difference in slope rather than an offset in the scatter plot.

The good agreement in the pattern of fine structure of HCHO concentrations for portions of the campaign suggest that, at least for those periods, the instruments were both sampling the same air mass. Systematic errors in calibrations were ruled out during the campaign. The offset during these periods of excellent agreement in trends is not significantly different from other sampling periods. Subsequent experiments, performed during the following summer at NPL, suggest that for laboratory conditions, the instruments were in good agreement.

Obviously one, or both, instruments were subject to systematic errors during NAMBLEX, but either these had been rectified by the following summer, or are only apparent

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during real air, rather than laboratory sampling. We are therefore at a loss to explain the differences in measured concentrations at Mace Head during NAMBLEX and in the following section the results of each apparatus are presented separately. The measurements are compared with other measurements and various models in the discussion section.

3.2. Average values and diurnal profiles for UoL apparatus

The campaign can broadly be divided into three different air masses. From 1–5 August the site experience continental air masses which had originated in Scandinavia and had passed over northern England. This was followed by a long period (6–17 August) of relatively long trajectories (high wind speeds) with a significant westerly component (ranging from south to north westerly). It is possible to further subdivide this period into air masses with NW, W, and SW origins, however, analysis showed no significant variation in formaldehyde concentrations with these finer components and hence all westerlies are considered together. Over the last few days of the formaldehyde campaign the wind was still westerly but from an anticyclonic weather system with low wind speeds. Summaries of the formaldehyde concentrations for each period can be found in Table 2.

Not surprisingly, formaldehyde concentrations were highest during the initial continental air flow. Concentrations of hydrocarbons and CO are also elevated during this period. Thereafter, concentrations decreased and a diurnal pattern became more apparent, Fig. 5a, with concentrations during the westerly air flows varying from an early morning (03:00–05:30 GMT) minimum of ~850 pptv to a peak value of ~1050 pptv between 14:00–16:00 GMT (local time is GMT + 1). During the latter part of the anticyclonic westerly airflows, the amplitude of the diurnal profile appeared to increase. There were also concurrent increases in acetylene concentrations suggesting some local contamination during these periods.

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3.3. Average values and diurnal profiles for UEA apparatus

Results from the fluorescence apparatus show generally similar behaviour. Concentra-
tions are elevated at the start of the campaign, but appear to remain constant between
200–500 pptv from the 3–14 August. As with the UoL data for the westerly winds, a pro-
nounced diurnal variation is observed, as shown in Fig. 5b, with an amplitude similar
to that observed by the UoL.

The concentrations measured on the 15 August are significantly lower than other
periods during the westerly airflow with values between 50–100 pptv. After the 16
August data from the UEA apparatus became patchy due to intermittent lamp failure.

4. Discussion

4.1. Comparison with previous marine boundary layer measurements

Some results from previous campaigns are presented in Table 1. We have focused pre-
dominantly on measurements from the North Atlantic although some other campaigns
are included.

Formaldehyde measurements at Mace Head are relatively limited. A measurement
campaign was conducted at Mace Head under the EMEP programme, with formalde-
hyde being monitored from November to April using 2,4 DNPH cartridges with subse-
quent off-line analysis by HPLC (Solberg et al., 1996). Samples were collected twice
weekly, each measurement consisting of 8 h sampling centred around midday, with
subsequent off line analysis at NILU. Solberg et al. (1996) reported monthly averages
between 200–500 pptv over this winter period. Direct comparison with our summer
data cannot be made, however, the winter concentrations appear to correlate well with
those made at a Norwegian site (Birkenes), where summer concentrations reach al-
most 1 ppbv. Solberg et al. (1996) report annual variations in formaldehyde levels from
a number of rural sites around Europe and all show summer maxima.

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Formaldehyde was measured at Mace Head during the ACSOE 96 and 97 campaigns. A majority of the measurements were made using an earlier version of the fluorometric technique, and a more limited data set was also recorded using tuneable diode laser spectroscopy (TDLS) (Cardenas et al., 2000). Typical values for westerly air masses were 200–400 pptv from the fluorometric technique, whereas the TDLS produced higher, but scattered values with a range of 0–1000 pptv.

A number of formaldehyde measurements have been made during the North Atlantic Regional Experiment (NARE) campaigns. During NARE 1993, Tanner et al. (1996) measured a range of carbonyl compounds using derivatization via 2,4 DNPH/HPLC at Cherbourg Point, Nova Scotia. Typical values for clean air at 200–400 pptv are in good agreement with the UEA data. Measurements were only made every six hours, so no diurnal trends are reported. Tanner et al. (1996) report a relationship between formaldehyde and ethanal with the latter being approximately 50% of the formaldehyde concentrations. This observation will be discussed further below.

During NARE 1997 a number of airborne measurements were made from St John's, Newfoundland, using both TDLS and the coil/DNPH technique (Fried et al., 2002). Generally the agreement between the two techniques was good with a gradient on the scatter plot of 0.98 ± 0.07 , however, approximately 30% of the data points lay outside the combined 2σ uncertainties. Median values in clean background air from 0–2 km, between 35–55° N are 400 pptv for TDLS (range 300–900 pptv for lowest altitude measurements) and 410 pptv for the CDNPH (range 200–800 pptv for lowest altitude measurements). Average concentrations decrease with both altitude and latitude.

Two shipborne campaigns have monitored formaldehyde in the North Atlantic. The ALBATROSS campaign took place during October and November 1996 with the cruise being a meridional track from 60° N to 45° S. Formaldehyde was monitored via a commercial apparatus based on the Hantzsch reaction (Weller et al., 2000). A variety of different air masses were encountered. Typical concentrations when the ship was between 20–40° N and intercepting trajectories similar to the westerlies and south-westerlies reaching Mace Head were in the region of 300–800 pptv. Harris et al. (1992)

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report data from the 1988 Polar Stern cruise which took place during September and October from 45° N into the southern hemisphere. Formaldehyde was predominantly measured using TDLS with a small number of comparative experiments obtained with DNPH sampling and subsequent analysis via HPLC. The two methods were in good agreement. Harris et al report a mean formaldehyde concentration between 40–45° N of 650 pptv, with concentrations decreasing slightly during the cruise south.

When used to compare with formaldehyde concentrations at Mace Head, the results of other campaigns need to be interpreted carefully. For air and shipborne measurements variation with latitude will be strongly dependent on the origins of the air mass reaching the receptor at that point and may not be typical of longer term measurements at that latitude. Both Fried et al. (2002) and Weller et al. (2000) report a decrease in average formaldehyde concentrations with latitude, however, because the air masses trajectories and ocean currents vary across the region, this does not mean that the latitudinal variation will be the same over the eastern Atlantic. Finally, the EMEP data showed significant annual variation in formaldehyde concentrations (as would be expected from a photochemical intermediate) and therefore results from different seasons should be interpreted with care.

The final campaign discussed in this section took place in the Indian Ocean during spring 1999 (Wagner et al., 2001). Given the location, comparison with Mace Head concentrations are difficult, however, results from the INDOEX campaign, (where formaldehyde was measured by TDLS) are interesting for several reasons. Firstly, even for trajectories that had been over the open ocean for seven days, and hence lost any continental formaldehyde, concentrations were still significant (~500 pptv). A direct oceanic source is not expected and therefore this is good evidence of significant HCHO production from long-lived precursors, including oxygenated VOC. Secondly, a strong positive correlation was noted with CO for a variety of air masses including very aged air ($\text{HCHO/CO} \approx 3 \times 10^{-3}$). Finally, during periods of consistent air flow, diurnal variations in [HCHO] were observed with an amplitude of approximately 200 pptv in the northern hemisphere, consistent with the observed amplitudes of both UEA and UoL

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instruments during clean westerly airflows.

4.2. Comparison with other NAMBLEX measurements

The NAMBLEX campaign was characterised by relatively polluted air during the first part of the campaign, with a period of consistent, strong, westerly, clean air and finally anticyclonic conditions, originating in the North Atlantic. Anthropogenic pollutants are therefore expected to be high during the initial part of the campaign, and thereafter relatively low, although the low wind speeds under anticyclonic conditions may allow for influence by local sources. The acetylene and CO concentrations shown in Fig. 6 broadly match the expected behaviour.

If the correlation between formaldehyde and CO, observed during the INDOEX campaign by Wagner et al. (2001) applied under NAMBLEX conditions, then formaldehyde concentrations during westerly airflows would be predicted to be in the region of 200–300 pptv, in good agreement with the UEA data. However, Fig. 6 shows a strong correlation between CO and acetylene, a primary pollutant. Therefore CO concentrations at Mace Head may correlate more strongly with primary sources rather than photochemical production and the relationship between formaldehyde and CO observed during highly aged air in the Indian Ocean may not apply.

Methane levels are likely to be slightly elevated in polluted air masses, but formaldehyde is also formed during the atmospheric oxidation of virtually all higher hydrocarbons and oxygenated VOCs (OVOC), and due to their higher photochemical loss rates, these latter species are likely to be the dominant influence on formaldehyde concentrations.

A good example of a formaldehyde precursor is acetaldehyde (ethanal). Both photolysis and reaction with OH can lead to formaldehyde formation:



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Figure 7a shows a comparison of the time series for acetaldehyde and formaldehyde, with Fig. 7b showing the corresponding correlation plots with the UoL data. Acetaldehyde was measured by the University of York (Hopkins et al., 2003; Lewis et al., 2005) concentrating VOC and OVOC from ~1 litre of air onto a solid trap followed by rapid thermal desorption, separation via GC with detection by FID. The general correlation between the CH₃CHO and H₂CO data from the UoL is good with both showing the same gradual decline in concentrations at the beginning of the campaign, followed by consistent and relatively low levels during the periods of westerly air flow.

The correlation plot of acetaldehyde concentrations with the UoL formaldehyde data shows an intercept of 0.46 ppbv on the HCHO axis. An intercept would be expected given that a significant fraction of formaldehyde will be formed from methane or from fragmentation processes that by-pass acetaldehyde production.

A significant positive correlation between formaldehyde and acetaldehyde was observed by Tanner et al. (1996) during measurements at a coastal site in Nova Scotia during NARE 1993. They reported $[\text{HCHO}]_{\text{ppbv}} = 1.9[\text{CH}_3\text{CHO}]_{\text{ppbv}} + 0.22$ with an *r* coefficient of 0.62. Errors on the gradient and intercept are not reported, but from the scatter of the plot, are likely to be at least ±33% and therefore comparable with our parameters derived observations between acetaldehyde and the UoL formaldehyde data. Monthly averaged formaldehyde and acetaldehyde concentrations were reported by Solberg et al. (1996) from a number of remote European sites including Mace Head. In all cases formaldehyde concentrations were greater than acetaldehyde, typically about a factor of two during the summer months.

The UEA formaldehyde concentrations are comparable in value with the measured acetaldehyde concentrations. Given our current understanding of formaldehyde and acetaldehyde photo oxidation these results appear to be incompatible.

Methanol is another significant OVOC precursor for formaldehyde. Reaction with OH leads to HCHO via reaction of either CH₃O isomer formed by the initial abstraction reaction. In the MBL reaction of Cl atoms with hydrocarbons and OVOCs may also

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need to be considered (Ramacher et al., 1999). Cl initiated reactions tend to be less selective in the position of abstraction, but in general form the same type of radicals as OH initiation. Although concentrations of Cl atoms are much lower than OH in the MBL (typically $5 \times 10^3 \text{ cm}^{-3}$ vs. $1 \times 10^6 \text{ cm}^{-3}$), the magnitude of the abstraction rate coefficients (typically $>5 \times 10^{-11} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$; Qian et al., 2002; Seakins et al., 2004) is such that Cl initiated chemistry can be significant.

4.3. Modelling

The simplicity of the chemistry in the MBL allows for the construction of relatively simple models to test our understanding of photochemistry in this environment. The ideal molecules for comparison are those with very short lifetimes, so that transport can be ignored and realistic comparisons can be made with box models. Hydroxyl and hydroperoxy radicals are ideal examples and comparisons between model predictions and measurements have been made in a number of environments, including the marine boundary layer (Carslaw et al., 2002; Heard et al., 2004; Sommariva et al., 2004). The lifetime of formaldehyde (typically $\sim 4 \text{ h}$ under midday conditions) is such that transport should really be considered, however, zero dimensional box models give an indication of the expected concentrations from various chemical schemes and highlight the relative importance of various formation and removal channels. A good example of a box model is described by Wagner et al. (2001, 2002) and used to interpret shipborne measurements in the remote Indian Ocean from the INDOEX campaign. In this case methane was the dominant source of formaldehyde and good agreement (typically within 20%) was found between measurements and modelled formaldehyde concentrations. However, as shown in Table 3, in other campaigns the models can either under or over predict formaldehyde concentrations.

In this study we have used the Master Chemical Mechanism (MCM) (Saunders et al., 2003) to simulate the formaldehyde concentration based on two models. The 'clean' model used only a very small subset of the MCM based on methane and CO chemistry whilst the 'full-oxy' model was based on a mechanism originating from CO, CH₄,

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23 hydrocarbons (including isoprene) (Lewis et al., 2005), DMS, chloroform and three oxygenated compounds (methanol, acetaldehyde and acetone) (Hopkins et al., 2003; Lewis et al., 2005). During the simulations the model was constrained to 15 min averages of the hydrocarbons, H_2 , O_3 , NO , NO_2 , H_2 , temperature and the measured photolysis rates of O_3 , NO_2 , $HONO$, $HCHO$, CH_3COCH_3 and CH_3CHO . The model included parameters for heterogeneous uptake and dry deposition and was run for several days (typically 2–3) to initialize the values of some species which were not constrained.

Figures 8a and b show comparisons of the models with the experimental data for two periods. Figure 8a is for JD221–222 during the westerly airflows when both instruments observed a pronounced diurnal variation. For these conditions the full-oxy model is clearly in better agreement with the UEA data, although the shape of the diurnal pattern does not produce the diurnal profiles observed by either technique, particularly the decrease in concentration in the afternoon and evening of JD221.

Figure 8b shows a comparison of the clean and full-oxy models with the experimental data for the period 15–21 August. The predictions from the full-oxy model are in reasonable agreement with the formaldehyde levels observed from UoL apparatus. In general the model does not do a particularly good job in predicting the fine structure of the formaldehyde concentrations (an exception being the sharp increase in concentrations in the early afternoon of 16 August, observed by both experimental techniques), but this is to be expected from a box model.

Figure 8b also highlights the significant contribution that the higher hydrocarbons and particularly the oxygenated species make to the predicted formaldehyde concentration. This observation is in good agreement with the analysis by Wagner (2002), where even in the much less complex conditions of the remote Indian Ocean, only 77% of $HCHO$ originates from methane, the other 23% coming from ethane, ethene, propene, isoprene, acetone and DMS, all at concentrations significantly lower than those encountered at Mace Head. Our model highlights the importance of acetaldehyde on formaldehyde generation, but unfortunately acetaldehyde was not considered

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by Wagner.

A rate of production analysis also emphasises the domination (>70%) of the $\text{CH}_3\text{O}_2 + \text{NO}$ reaction as the loss mechanism for CH_3O_2 for $[\text{NO}]$ in the region of 10 pptv. Under these conditions formation of the CH_3OOH reservoir is limited and therefore the potential for carbon removal via CH_3OOH deposition is relatively small.

There have been several recent formaldehyde measurement/model comparisons in the North Atlantic environment (Weller et al., 2000; Frost et al., 2002; Fried et al., 2003) which all under predict measured formaldehyde concentrations to some degree. Weller et al. (2000) compared measured average formaldehyde concentrations of 580 ± 160 pptv from a North Atlantic cruise with calculated values based on a box model. On the basis of measured alkane concentrations, they argued that methane should be the major source, but noted that alkenes (not measured) could be a significant source. The standard methane model under predicted the measurements by 250 pptv. Better agreement was found by introducing a formaldehyde production channel from Reaction (R2);



based on arguments by Ayers et al. (1997). Making channel (R17) 40% of the total Reaction (R2) increased average formaldehyde concentrations by 80 pptv. Increasing the rate coefficient for OH abstraction and changing the branching ratios for the OH + methylhydrogenperoxide reaction within values suggested in the literature increased formaldehyde by 50 pptv over the base model. Frost et al. (2002) generally underestimated airborne formaldehyde measurements from the NARE 97 campaign by between 130 and 180 pptv (~ factor two) using a methane/hydrocarbon model. They carried out a sensitivity analysis and concluded that it was not possible to bridge the gap between measurement and model within the currently understood uncertainties in the model parameters but noted that VOCs not measured or considered in the model could represent the missing source. Finally, Fried et al. (2003) report generally good agreement between measurement and model for the TOPSE 2000 experiment, although again

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there is significant under prediction at high latitudes over North Atlantic regions (Fried et al., 2003).

Our modelling work emphasises the importance of oxygenated species and particularly acetaldehyde and methanol in generating significant concentrations of formaldehyde. We would therefore suggest that these species, measured values of which were not considered in the above modelling studies, as a potential source of missing formaldehyde and recommend that concurrent measurements of all oxygenated species are considered for future campaigns.

During the night (from 20:00–04:00 h) deposition and entrainment are the only removal processes for formaldehyde. From the averaged diurnal formaldehyde profiles for westerly air masses (Figs. 5a, b) both experimental techniques show a decrease of approximately 150 pptv formaldehyde during this period. Assuming that physical processes can be represented by a first order loss, this allows a prediction of the initial concentration at the start of the deposition only period.

$$[\text{HCHO}]_0 - 150 = [\text{HCHO}]_0 e^{-kt} \quad (1)$$

For a sum of deposition and entrainment equal to $1 \times 10^{-5} \text{ s}^{-1}$, based on the reported deposition and entrainment velocities of 0.4 cm s^{-1} (Wagner et al., 2002) and a boundary layer height of 800 m, Eq. (1) predicts $[\text{HCHO}]_0$ of 600 pptv, in between the two observations. The uncertainties in boundary layer height and deposition velocities are such that the calculation range can encompass both observations. However, reductions in such uncertainties could help differentiate between the two observations.

4.4. Implications for HO_x production

Formaldehyde is an important photolytic source of radicals in the troposphere (R10b). The MCM has been used to quantify the influence of the measured formaldehyde levels on $[\text{HO}_x]$. Variations in formaldehyde levels have relatively little influence on OH radical concentrations as while increased formaldehyde generates more HO_x radicals, formaldehyde is also a significant HO sink via Reaction (R8). Using either set of

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formaldehyde measurements changed the predicted OH concentration by <2% under typical westerly conditions. Increased formaldehyde concentrations have a more pronounced positive effect on HO₂ chemistry, although the relationship is not 1:1, being buffered by relatively complex radical interconversion. Increasing formaldehyde concentrations by 50% produces between a 15–25% increase in HO₂ concentrations. It should be noted that currently (Sommariva 2004) the MCM tends to over-predict HO₂ concentrations by up to a factor of two. As noted earlier, formaldehyde may have a more pronounced influence on HO_x production in other environments e.g. where a lack of ozone or water prevents OH generation via O(¹D) production.

5. Conclusions

Formaldehyde concentrations, measured by two different techniques during the NAMBLEX campaign appear to be correlated, but exhibit a significant offset. Joint calibrations and subsequent intercomparison experiments have failed to identify the cause of the systematic errors present in one, or both sets of apparatus. Formaldehyde continues to be a difficult molecule to measure. A recent comparison from the BERLIOZ campaign (Grossmann et al., 2003) reported formaldehyde measurements from a commercial Hantzsch system to be below that recorded by DOAS, however, Grossman reports that the DOAS recorded higher values in an intercomparison in the EUPHORE chamber. Clearly more work is required to identify systematic errors in formaldehyde detection.

Comparison with previous formaldehyde measurements in the North Atlantic suggests typical concentrations between 300–800 pptv under clean background conditions. The UEA data lie at the low end of this range, the UoL data just above the upper limit.

Previous campaigns have shown a strong positive correlation between formaldehyde and acetaldehyde and with a HCHO to CH₃HCO ratio of approximately 2:1. A good correlation has been observed between the UoL formaldehyde data and acetaldehyde

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measurements from the University of York. The ratio of concentrations is in agreement with previous work.

Both instruments show a distinct diurnal profile during periods of consistent westerly airflow with an amplitude of 150–250 pptv. This is the expected behaviour of a photochemical intermediate, but such behaviour has not always been observed in previous campaigns, either due to sampling frequency or because of fluctuations in the concentrations of the various air masses arriving at the receptor. This latter factor is especially important in shipborne cruises.

Modelling studies from the latter part of the campaign predict concentrations between the two experimental measurements, but more importantly, highlight the importance of oxygenates in formaldehyde production. The involvement of oxygenated species in formaldehyde production may explain some of the under predictions of simple methane only models. The concentrations of formaldehyde predicted by the MCM for an earlier part of the campaign are in better agreement with UEA data.

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Table 1. Some previous measurements of formaldehyde in the marine environment (predominantly North Atlantic).

Reference	Campaign and Date	Location	Platform	Technique	[HCHO]/pptv	Notes
Harris et al., 1992	Polarstern 1988	North Atlantic	Ship	TDLS	650	Measurement is mean value over 40–45° N. No diurnal variation.
Tanner et al., 1996	NARE 1993	Nova Scotia	Coastal site	DNPH	200–400	Summer measurement
Solberg et al., 1996	EMEP 1994–1995	Mace Head	Coastal site	DNPH	200–500	Winter measurement. Measurements at other sites show summer maxima.
Cardenas et al., 2000	ACSOE 1996	Mace Head	Coastal site	Hantzsch	200–450	Summer measurement.
Weller et al., 2000	ALBATROSS 1996	North Atlantic	Ship	Hantzsch	400–500	No diurnal variation observed. Autumn campaign.
Fried et al., 2002	NARE 1997	Newfoundland	Plane	TDLS	410±150	
Wagner et al., 2001	INDOEX 1999	Indian Ocean	Ship	TDLS	430±100	Diurnal variation with amplitude of ~200 pptv observed.

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Table 2. Summary of UoL and UEA formaldehyde concentrations for the various air masses encountered during NAMBLEX. All concentrations in ppb.

Air mass origin	UoL		UEA	
	24 h average [HCHO]	Range	24 h average [HCHO]	Range
Continental	1.20±0.13	0.98–1.58	0.37±0.11	0.13–0.78
Westerly	0.96±0.09	0.78–1.15	0.27±0.07	0.13–0.43
Anticyclonic	0.92±0.15	0.72–1.31		

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Table 3. Summary of the results from some previous measurement/model intercomparisons.

Authors	Environment	Model constraints	Typical [HCHO]/pptv	Under/Over prediction
Weller et al., 2000	Marine, Atlantic, 45° N–35° S	Photochemical box model based on CH ₄ and CO photo-oxidation	580±160 ^a	Under predicted by factor 2
Ayers et al., 1997	Coastal, Cape Grim, Tasmania	Photochemical box model based on CH ₄ and CO, and including DMS	400±50 ^b	Under predicted by factor 2
Wagner et al., 2002	Marine, Indian Ocean	MCM 2 Photochemical oxidation of CH ₄ and selected NMVOC.	200±70 ^a	Over predicted by 12%
Zhou et al., 1996	Free tropospheric air	Simple box model based on CH ₄ , constrained to measured CH ₃ OOH	211±104 ^c	Over predict
Liu et al., 1992	Free tropospheric air	Photochemical box model based on CH ₄	105±42	Over predicted
Jacob et al., 1996	South Atlantic (Aircraft)	Photochemical box model includes CH ₄ and NMVOCs, but not CH ₃ OH and CH ₃ CHO.	110	Over predicted
Fried et al., 2003	Marine, Pacific (Aircraft)	NASA Langley box model, constrained by observations including oVOCs.	(Average 2–4 km) 196±161	Equivalent
Frost et al., 2002	Marine, North Atlantic (Aircraft)	Photochemical box model includes NMHC and constrained to measurements (some, typical values from other campaigns).	(Average 0–2 km) 410±110	Under predicted by a factor of 2

^a Daily mean^b Value read from graph^a Summer intensive

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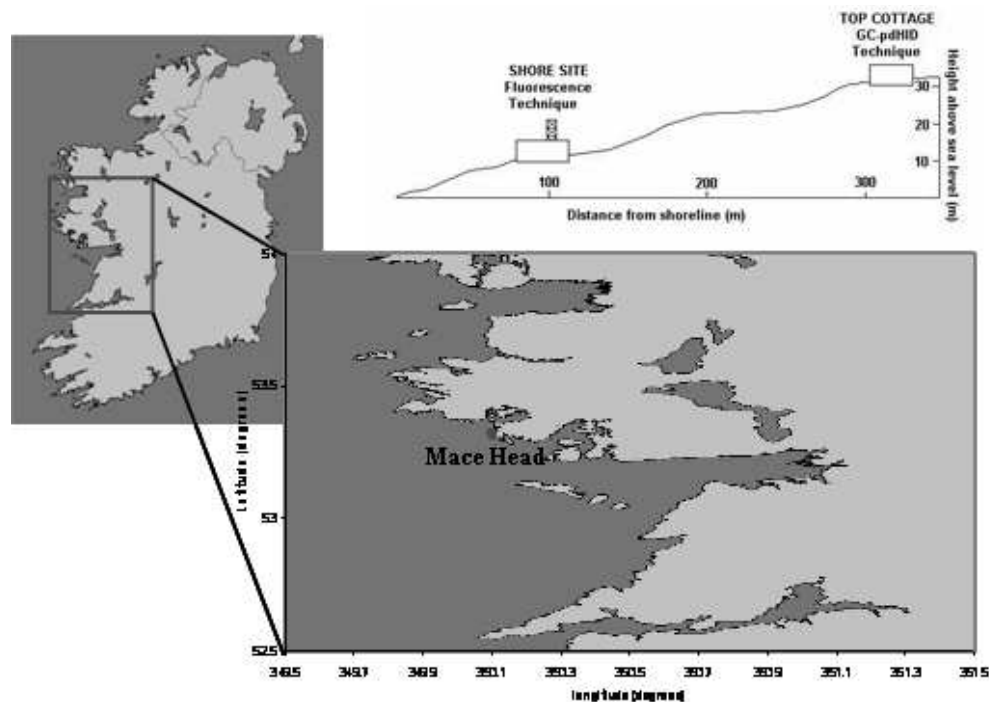


Fig. 1. Location of the Mace Head observatory and the relative positioning of the two formaldehyde instruments.

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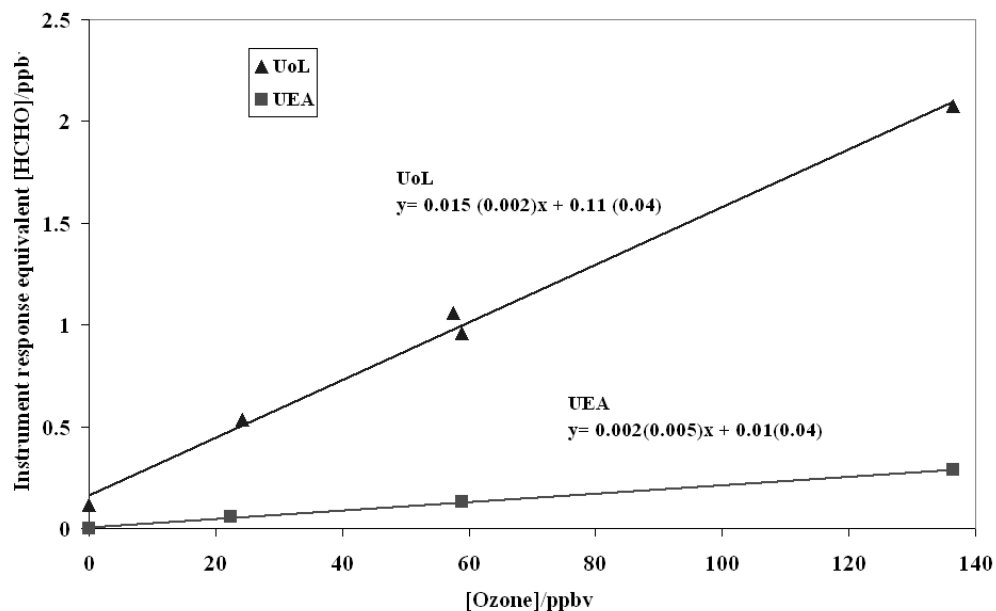


Fig. 2. Ozone artefact for (Δ) UoL apparatus, (\square) UEA apparatus as determined during experiments at NPL, 2003.

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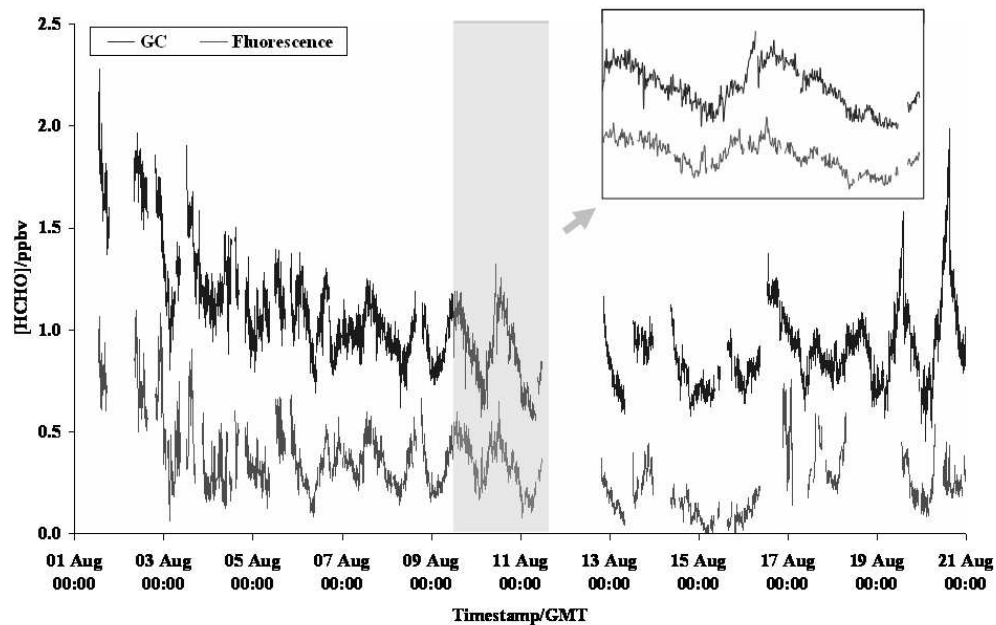


Fig. 3. Time series of formaldehyde concentrations from GC and fluorescence techniques. All data corrected for ozone interferences.

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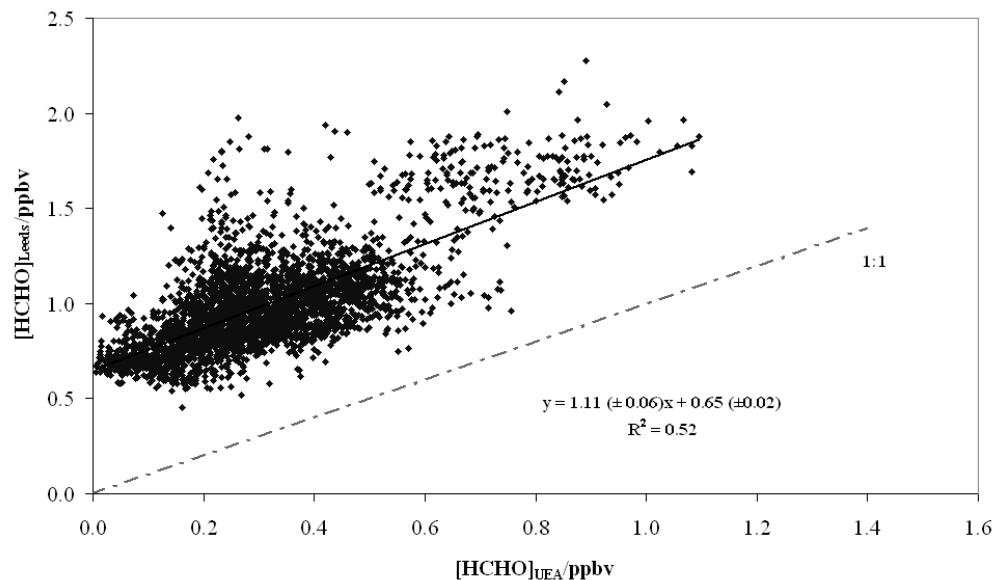


Fig. 4. A plot showing the weighted bivariate regression between the GC and the fluorescence technique for all data (solid line) and the 1:1 correlation (dotted line).

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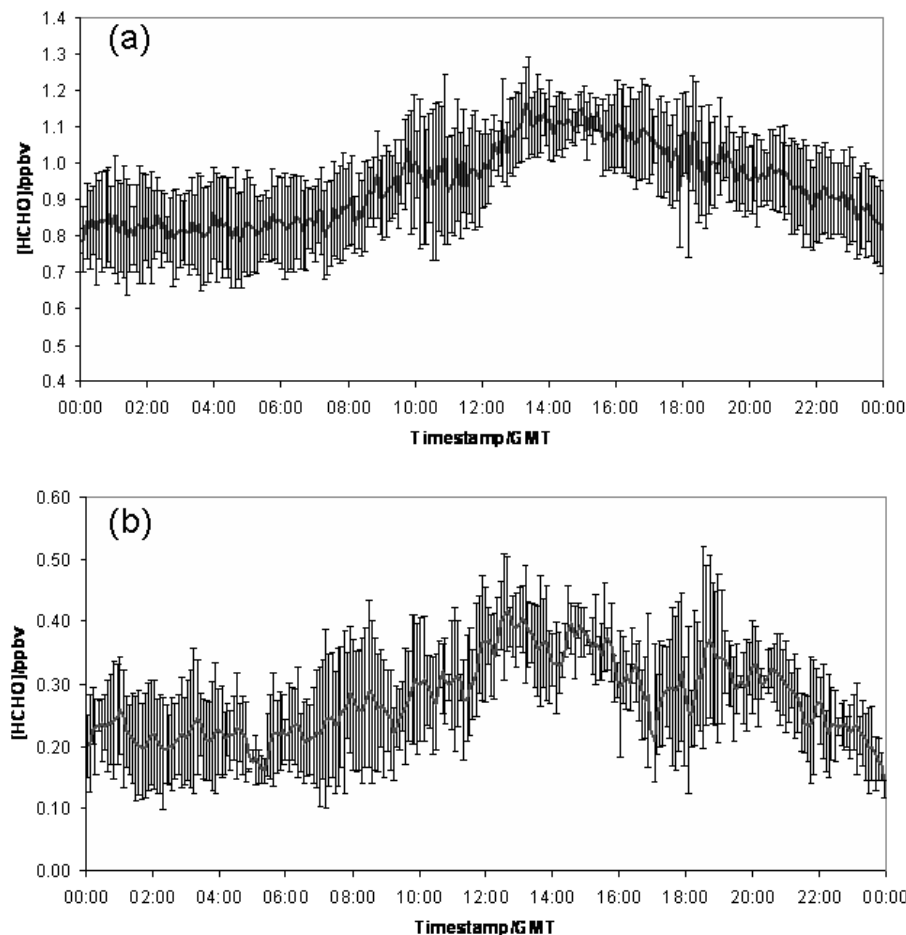


Fig. 5. Diurnal variation of formaldehyde concentrations measured by (a) UoL, (b) UEA during westerly airflows (6–17 August 2002). Error bars represent one standard deviation.

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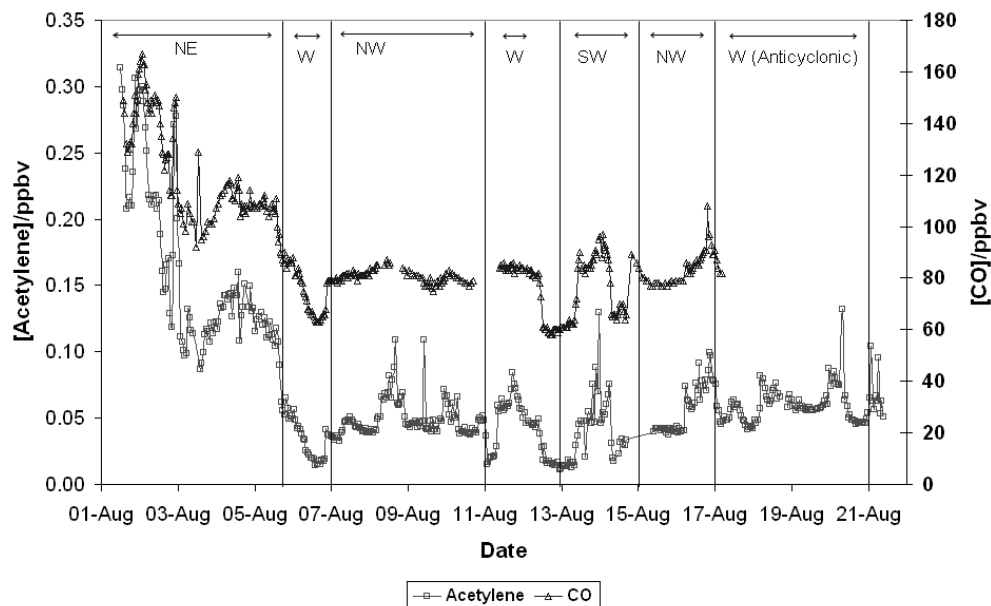


Fig. 6. Signatures of anthropogenic tracers. Acetylene and CO data were provided by the University of York and Mace Head Research Station, respectively.

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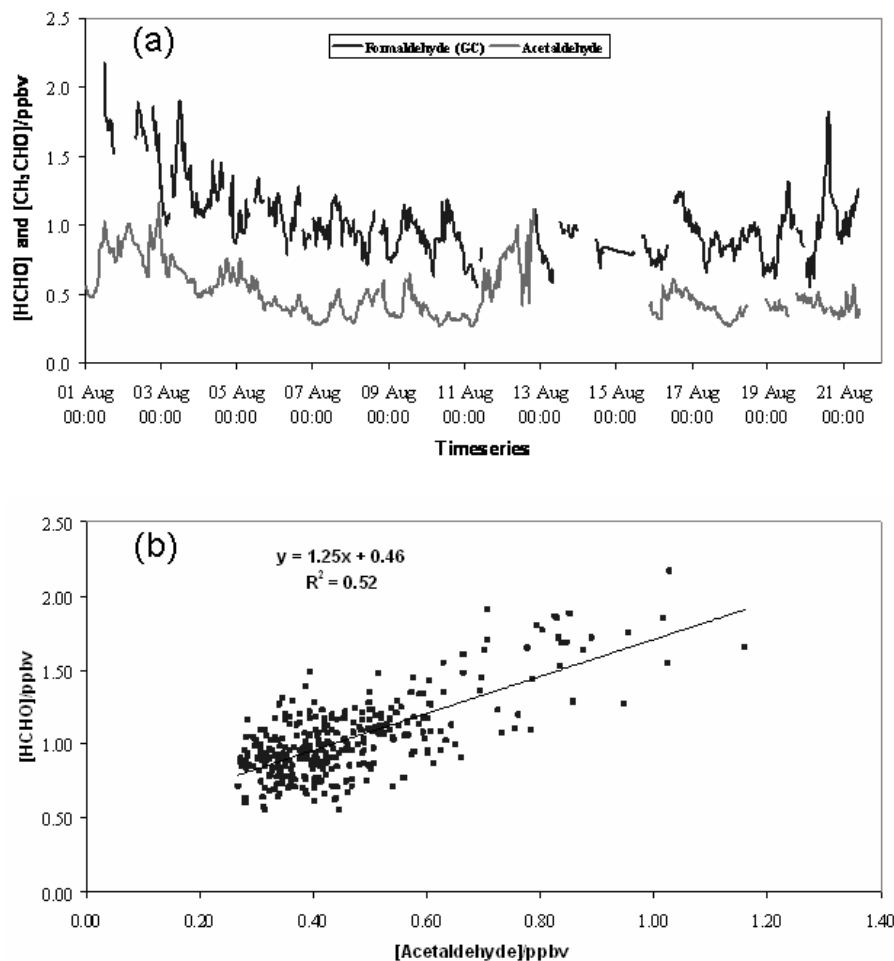


Fig. 7. (a) Time matched series of formaldehyde (UoL) and acetaldehyde (University of York) concentrations. (b) Correlation plot of the formaldehyde and acetaldehyde concentration data.

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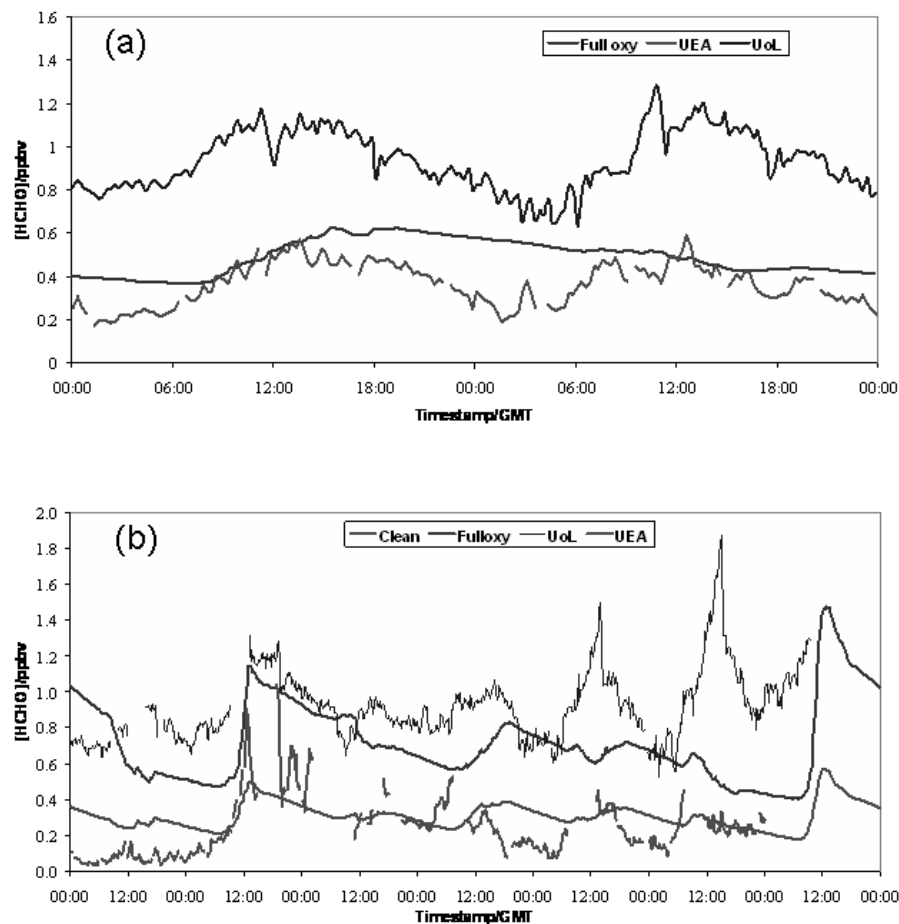


Fig. 8. (a) Comparison of measured and modelled formaldehyde concentrations for 9–10 August 2002 (JD221–222). (b) Comparison of measured and modelled formaldehyde concentrations for 15–21 August 2002 (JD227–233).

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